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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09/900,647	07/07/2001	Dale R. Lovercheck	ANAL-VIT	6584	
75	90 07/02/2003				
Dale R. Lovercheck, Esquire			EXAMINER HUI, SAN MING R		
92 Patricia Place Media, PA 19063					
			ART UNIT	PAPER NUMBER	
		•	1617	1<	
			DATE MAILED: 07/02/2003	•	

Please find below and/or attached an Office communication concerning this application or proceeding.

•		Application No.	Applicant(s)	
•		09/900,647	LOVERCHECK, DALE R.	•
Office Action S	ummary	Examiner	Art Unit	
•		San-ming Hui	1617	
	this communication a		eet with the correspondence address -	
 If NO period for reply is specified above Failure to reply within the set or extend Any reply received by the Office later to earned patent term adjustment. See 3 	IS COMMUNICATION noter the provisions of 37 CFR g date of this communication. Is less than thirty (30) days, a re, the maximum statutory perioded period for reply will, by state han three months after the maximum stater s	N. 1.136(a). In no event, however, reply within the statutory minimur od will apply and will expire SIX of tute, cause the application to be	may a reply be timely filed n of thirty (30) days will be considered timely. 6) MONTHS from the mailing date of this communica ome ABANDONED (35 U.S.C. § 133).	ition.
Status 1) Popposity to community	unication(s) filed on 1	4 March 2002		
1) Responsive to commu	· · · —	<u> </u>		
2a) This action is FINAL .	,—	This action is non-final.		
			al matters, prosecution as to the merit 35 C.D. 11, 453 O.G. 213.	S IS
4) Claim(s) 26-30,33-35	and 37-46 is/are pend	ding in the application.	•	
4a) Of the above claim((s) is/are withd	rawn from consideratio	n.	
5) Claim(s) is/are a	allowed.			
6)⊠ Claim(s) <u>26-30, 33-35,</u>	and 37-46 is/are reje	ected.		
7) Claim(s) is/are o	objected to.			
8) Claim(s) are sub	oject to restriction and	d/or election requireme	nt.	
Application Papers	•			
9) ☐ The specification is object	ected to by the Exami	ner.		
10) The drawing(s) filed on	is/are: a)□ ac	cepted or b) objected t	by the Examiner.	
Applicant may not reque	est that any objection to	the drawing(s) be held in	abeyance. See 37 CFR 1.85(a).	•
11) ☐ The proposed drawing of	correction filed on	is: a)⊡ approved b) disapproved by the Examiner.	
	-	reply to this Office action		
12) The oath or declaration	•	Examiner.		
Pri rity under 35 U.S.C. §§ 119	and 120		·	
13) Acknowledgment is ma	ade of a claim for fore	ign priority under 35 U.	S.C. § 119(a)-(d) or (f).	
a)□ All b)□ Some * c)[☐ None of:			•
1. Certified copies	of the priority docume	ents have been received	i.	
2. Certified copies	of the priority docume	ents have been received	d in Application No	
	om the International E	Bureau (PCT Rule 17.2		
14) Acknowledgment is mad	e of a claim for dome	stic priority under 35 U	S.C. § 119(e) (to a provisional application	ation).
a) ☐ The translation of t 15)☐ Acknowledgment is mad	he foreign language p	provisional application I	nas been received.	ŕ
Attachment(s)		•		
1) Notice of References Cited (PTO-8 2) Notice of Draftsperson's Patent Dr. 3) Information Disclosure Statement(s	awing Review (PTO-948)	5) 🔲 Not	rview Summary (PTO-413) Paper No(s)ice of Informal Patent Application (PTO-152) er:	
S. Patent and Trademark Office PTO-326 (Rev. 04-01)	Office	Action Summary	Part of Paper No. 15	

Art Unit: 1617

DETAILED ACTION

Upon reconsideration, the finality of the rejection of the last Office action is withdrawn.

Claims 26-30, 33-35, and 37-46 are pending.

The elected species are ibuprofen as the discomfort reliever and vitamin C as the nutritional supplement. The election was made in Paper No. 5.

Claim Rejections - 35 USC § 112

Claims 26, 29-30, 33-35, 37-38, 40-44, and 46 are rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the enablement requirement. The claim(s) contains subject matter which was not described in the specification in such a way as to enable one skilled in the art to which it pertains, or with which it is most nearly connected, to make and/or use the invention. The instant specification does not provide sufficient information for one skilled artisan to practice the instant invention without undue experimentation. The instant claims encompass "nutritional supplement not being adapted to aid in or contribute to discomfort relieving of said discomfort reliever, said nutritional supplemental not being adapted to aid in or contribute to reducing side effects of said discomfort reliever". The instant claims also recite the preferred nutritional supplement as vitamin C (See claim 27). However, based on the teachings of the abstract of Tsunoda (reference of record), vitamin C is working synergistically with ibuprofen in pain relief (See the abstract). It is clear from the evidence of Tsunoda that the herein preferred recited nutritional supplement, vitamin C, actually contributes

to discomfort relieving of herein preferred recited discomfort reliever, ibuprofen. It is not clear how the recited method is enabled in view of the evidence of Tsunoda. Thus, the instant claims fails to comply with 35 USC 112, first paragraph.

Claims 26, 29-30, 33-35, 37-38, 40-44, and 46 are rejected under 35 U.S.C. 112, first paragraph, because the specification, while being enabling for the discomfort reliever recited in claim 32, does not reasonably provide enablement for other discomfort reliever suitable for the instant invention. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to use the invention commensurate in scope with these claims. In the instant case, the specification fails to provide sufficient information for enabling one skilled of artisan to practice the instant invention undue experimentation. Attention is directed to *In re Wands*, 8 USPQ2d 1400 (CAFC 1988) at 1404 where the court set forth the eight factors to consider when assessing if a disclosure would have required undue experimentation. Citing *Ex parte Forman*, 230 USPQ 546 (BdApls 1986) at 547 the court recited eight factors:

- 1) the quantity of experimentation necessary,
- 2) the amount of direction or guidance provided,
- 3) the presence of absence of working examples,
- 4) the nature of the invention,
- 5) the state of the prior art,
- 6) the relative skill of those in the art

Art Unit: 1617

- 7) the predictability of the art, and
- 8) the breadth of the claims.

Applicant fails to set forth the criteria that define "a discomfort reliever not being adapted to aid in or contribute to nutrition supplementing of said nutritional supplement". It is not clear what compounds would possess such characteristics. The instant specification fails to provide sufficient guidance to ascertain such characteristics. Additionally, Applicant fails to provide information allowing the skilled artisan to ascertain these compounds without undue experimentation. In the instant case, only a limited number of "a discomfort reliever not being adapted to aid in or contribute to nutrition supplementing of said nutritional supplement" examples are set forth, thereby failing to provide sufficient working examples. It is noted that these examples are neither exhaustive, nor define the class of compounds required. The pharmaceutical art is unpredictable, requiring each embodiment to be individually assessed for physiological activity. The instant claims read on all "a discomfort reliever not being adapted to aid in or contribute to nutrition supplementing of said nutritional supplement". necessitating an exhaustive search for the embodiments suitable to practice the claimed invention. Applicants fail to provide information sufficient to practice the claimed invention, absent undue experimentation.

Furthermore, Applicant uses functional language in attempt to define the instant invention. Attention is directed to *General Electric Company v. Wabash Appliance*Corporation et al 37 USPQ 466 (US 1938), at 469, speaking to functional language at the point of novelty as herein employed: "the vice of a functional claim exists not only

Art Unit: 1617

when a claims is "wholly" functional, if that is ever true, but when the inventor is painstaking when he recites what has already been seen, and then uses conveniently functional language at the exact point of novelty". Functional language at the point of novelty, as herein employed by Applicants, is further admonished in *University of* California v. Eli Lilly and Co. 43 USPQ2d 1398 (CAFC 1997) at 1406: stating this usage does "little more than outlin[e] goals appellants hope the recited invention achieves and the problems the invention will hopefully ameliorate". Applicants functional language at the point of novelty fails to meet the requirements set forth under 35 USC 112, first paragraph. Claims employing functional language at the point of novelty, such as Applicants', neither provide those elements required to practice the inventions, nor "inform the public during the life of the patent of the limits of the monopoly asserted" General Electric Company v. Wabash Appliance Corporation et supra, at 468. Claims thus constructed provide no guidance as to medicaments employed, levels for providing therapeutic benefit, or provide notice for those practicing in the art, limits of protection. Simply stated, the presented claims are an invitation to experiment, not reciting a specific medicament regimen useful for practicing the instant invention.

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 26-30, 33-35, and 37-46 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Art Unit: 1617

The expression "minor aches and pain <u>associated with</u> a common cold, ..., fatigue and drowsiness" recited in claims 26, 38, and 44 renders the claims indefinite as to the association recited herein. It is not clear what relationships between pain and other conditions are encompassed by the claims.

There are four "indicating" steps recited in claim 26. It is not clear how such indicating steps be accomplished. For example, it is not clear how the amount of the discomfort reliever be indicated? By imprinted onto the tablet? By information printed in the label? In the patient information leaflet? Engraved the amount onto the container? Thus, the metes and bounds of the claims cannot be ascertained.

The term "other upper respiratory allergy" recited in claims 26, 38, and 44 renders the claims indefinite as to the upper respiratory allergy encompassed by the claims.

The term "recommended daily value of nutritional supplement" in claims 38 and 44 renders the claims indefinite. As recited in claims 38 and 44, nutritional supplement can be herb (See claim 38, line 6 and claim 44, line 7). It is not clear what is the recommended daily value for herb, if such value exists. The instant specification does not define such term, and thus, the metes and bounds of the claims are not defined.

Where applicant acts as his or her own lexicographer to specifically define a term of a claim contrary to its ordinary meaning, the written description must clearly redefine the claim term and set forth the uncommon definition so as to put one reasonably skilled in the art on notice that the applicant intended to so redefine that claim term. *Process Control Corp. v. HydReclaim Corp.*, 190 F.3d 1350, 1357, 52 USPQ2d 1029, 1033 (Fed.

Art Unit: 1617

Cir. 1999). The term "indicator" in claims 29,33, 35, 38, 40, 42, and 44 is used by the claim to mean "a printed paper or a label" (See page 8, line 7-8), while the accepted meaning is:

- "1. One that indicates, as: a : an index hand (as on a dial) : pointer or Index b: an instrument, as a meter or gauge for monitoring the operation or condition of a physical system, as an engine, furnace, electrical network, or reservoir. c: The needle, dial, or other registering device on such an instrument.
- 2. Chem. A substance, as litmus or phenolphthalein, that indicates the presence, absence, or concentration of a substance or the degree of reaction between two or more substances by means of a characteristic change, esp. in color.
- 3. Any of various statistical values that collectively indicate the stability of an economic system."
- Webster's II New Riverside University Dictionary, 1984

The term is indefinite because the specification does not clearly redefine the term.

Claim 46 recites the limitation "discomfort is sleepiness, fatigue or drowsiness" in line 1. There is insufficient antecedent basis for this limitation in the claim. In claim 26, the discomfort is "minor aches or pain associated with ...". Sleepiness, fatigue, and drowsiness are not pain or minor aches.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

Art Unit: 1617

Claims 26-30, 33-35, and 37-46 are rejected under 35 U.S.C. 103(a) as being unpatentable over SS Pharmaceutical (Comline Biotechnology & Medical, 1 Dec. 1992, page 4), Tsunoda (JP 2000-229853, English abstract is also provided) and Yeh et al. (US Patent 5,032,384) in view of Krause (Krause's Food, Nutrition & Diet Therapy, 1992, page 277-279).

SS Pharmaceutical teaches a composition containing ibuprofen and a high content of vitamin C (See the abstract).

Tsunoda teaches a pain-alleviating tablet containing 300-500mg of ibuprofen and about 30-50mg of vitamin C (See the abstract).

Yeh et al. teaches a composition containing an antioxidant, such as ascorbic acid, and a NSAID, such as ibuprofen, such that the weight amount of the antioxidant and the NSAID is about 0.01 to 10% of the composition (See particularly the abstract, also col. 2, lines 9 and 48-49; col. 4, line 7-10). Yeh et al. also teaches that the composition can be formulated into oral dosage forms (See particularly col. 3, line 67).

The references do not expressly teach the composition to be indicated as in unit dosage form. The references do not expressly teach the composition to be indicated is in an enclosure. The references do not expressly teach the composition to be indicated is in an unit form as pill, tablet, or capsule. The references do not expressly teach the composition to be indicated is package with an indicator indicating the amount of each ingredients and the indication. The references do not expressly teach the indication of the recommended daily value of nutritional supplement.

Art Unit: 1617

Krause teaches that it is mandatory for nutrition manufacturer to list the recommended daily value of vitamin C of the food product on the package label (see page 279, Mandatory Listings of Food Label Section).

It would have been obvious to one of ordinary skill in the art at the time the invention was made to indicate the amount of ibuprofen and vitamin C, as a unit dosage form, in the composition claimed herein. It would have been obvious to one of ordinary skill in the art at the time the invention was made to enclose the ibuprofen-vitamin C unit dose tablet into a container with indicator (label) indicating the amount of each ingredients, the recommended daily value of the nutritional supplement, and the indication.

One of ordinary skill in the art would have been motivated to indicate the amount of ibuprofen and vitamin C, as a unit dosage form, in the composition claimed herein and enclose the same into a container with indicator (label) indicating the amount of each ingredients, the recommended daily value of the nutritional supplement, and the indication. Firstly, employing the herein claimed amount of ibuprofen and vitamin C is considered as optimization of result effect parameters, which is obvious as being within the skill of the artisan, absent evidence to the contrary. Secondly, putting the drug dosage form into a container is considered obvious within the purview of skilled artisan. Thirdly, inclusion of a package insert or label, which is considered as indicator in the instant case, showing the "the name of drug, dosage, dosage form, route of administration, indication and direction of use" of a pharmaceutical composition is mandated by 21 CFR 201.57 and is therefore obvious to one of ordinary skill in the art.

Art Unit: 1617

Finally, law also mandates listing the recommended daily value of a nutritional (Fig. Kraws, page >79, col. 1) supplement on the package label. Therefore, the method of indication of the herein claimed products is considered obvious to one of ordinary skill in the art since indicating the herein claimed information, regardless of what the drug is, is mandated by law.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to San-ming. Hui whose telephone number is (703) 305-1002. The examiner can normally be reached on Mon 9:00 to 1:00, Tu - Fri from 9:00 to 6:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Sreeni Padmanabhan, PhD., can be reached on (703) 305-1877. The fax phone numbers for the organization where this application or proceeding is assigned are (703) 308-4556 for regular communications and (703) 308-4556 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-1235.

San-ming Hui June 28, 2003 REENI PADMANABHAN PRIMARY EXAMINER